

HELLSTROM ET AL.
U.S. National Phase of PCT/EP2004/011958
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AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) A method of predicting whether an individual having hepatitis B virus (HBV) infection will respond to interferon (IFN) treatment; the method comprising; determining the presence or absence of antibodies reactive with a preS1 (94-117) peptide in a sample obtained from the individual,

the presence of said antibodies in said sample being indicative that said individual will respond to said treatment.

2. (Original) A method according to claim 1 comprising detecting the presence of said antibodies in said sample and thereby determining that the individual will respond to IFN treatment.

3. (Original) A method according to claim 1 comprising detecting the absence of said antibodies in said sample and thereby determining that the individual will not respond to IFN treatment.

4. (Currently Amended) A method according to ~~any one of the preceding~~ claim 1 wherein the individual has chronic HBV infection.

5. (Currently Amended) A method according to ~~any one of the preceding claims~~ claim 1 wherein the individual is HBeAg positive.
6. (Currently Amended) A method according to ~~any one of claims 1 to 4~~ claim 1 wherein the individual is HBeAg negative.
7. (Currently Amended) A method according to ~~any one of the preceding claims~~ claim 1 wherein the antibodies are IgG or IgM antibodies.
8. (Currently Amended) A method according to ~~any one of the preceding claims~~ claim 1 wherein the sample is a blood, serum or plasma sample.
9. (Currently Amended) A method according to ~~any one of the preceding claims~~ claim 1 comprising:
contacting the sample with a preS1 (94-117) peptide and;
determining binding of said antibodies to said peptide.
10. (Original) A method according to claim 9 wherein the peptide comprises a detectable label.
11. (Original) A method according to claim 9 wherein said peptide is immobilised.

12. (Currently Amended) A method according to ~~any one of claims~~claim 9 to 11 wherein said binding is detected with a labelled secondary antibody.

13. (Original) A kit for use in predicting whether an individual having hepatitis B will respond to interferon (IFN) treatment, the kit comprising:
a preS1 (94-117) peptide.

14. (Original) A kit according to claim 13 wherein said peptide is immobilised on a solid support.

15. (Original) A kit according to claim 14 wherein the solid support is a microtitre plate.

16. (Currently Amended) A kit according to ~~any one of claims~~claim 13 to 15 further comprising a labelled secondary antibody which binds to human antibodies.

17. (Currently Amended) A kit according to ~~any one of claims~~claim 13-16 further comprising regents for detecting the binding of the labelled secondary antibody.

18. (Currently Amended) A kit according to ~~any one of claims~~claim 13-17 further comprising wash buffers.

19. (Currently Amended) A kit according to ~~any one of claims~~claim 13-18 further comprising sample-handling containers.

20. (Currently Amended) A method of treating a hepatitis B infection in an individual comprising;

identifying the individual as responsive to interferon (IFN) treatment using a method according to ~~any one of claims~~claim 1-12, and;
administering IFN to said individual.

21. (Original) A method according to claim 20 wherein the IFN is alpha-IFN.

22. (Currently Amended) A method according to claim 20 or claim 21 wherein corticosteroid is administered to the individual.

23. (Currently Amended) A method of predicting whether an individual having hepatitis B virus (HBV) infection will respond to interferon (IFN) treatment which is substantially as described herein, ~~with reference to the accompanying table and figures.~~